

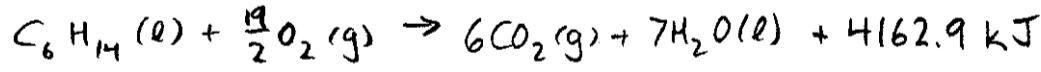
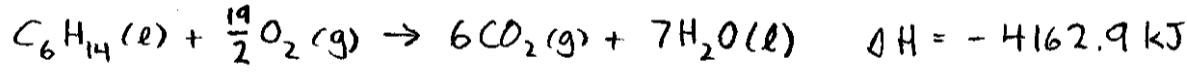
Name: _____

SCH OAI
Thermodynamics Test

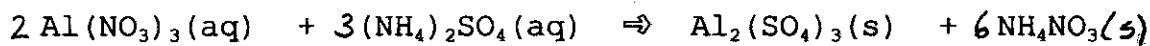
1. For each of the following, label with a + for an increase in potential energy or a - for a decrease in potential energy. Be careful. One-half mark will be deducted for each incorrect response.

- a) + The formation of ethyne from its component elements.
- b) - Sand is used to make glass.
- c) - Calcium carbonate is formed by the addition of carbon dioxide to calcium oxide.
- d) + Growth of a Western Red Cedar (i.e. humongous tree).
- e) - Cellular respiration.
- f) - Formation of a cloud.
- g) - Sublimation of $H_2O(g)$ to $H_2O(s)$.
- h) - Reaction of propane with oxygen.
- i) + Reduction of tin (IV) oxide to tin (II) oxide.
- j) + Conversion of glucose to hexane.

2. Please write the combustion reaction of hexane ($C_6H_{14}(l)$) as thermodynamic equation and as a thermochemical equation. The heat of combustion of hexane is -4162.9 kJ



3. Calculate the heat of reaction for:



Using the following heats of formation and heats of formation listed in your text.

$$\begin{aligned}\Delta H^\circ_{\text{Al}(\text{NO}_3)_3(\text{aq})} &= -273.65 \text{ kcal} \\ \Delta H^\circ_{(\text{NH}_4)_2\text{SO}_4(\text{aq})} &= -281.86 \text{ kcal} \\ \Delta H^\circ_{\text{Al}_2(\text{SO}_4)_3(\text{s})} &= -820.98 \text{ kcal}\end{aligned}$$

$$\begin{aligned}\Delta H^\circ_{\text{NH}_4\text{NO}_3(\text{s})} &= -365.6 \text{ kJ} \\ &= -87.38 \text{ kcal}\end{aligned}$$

Please note that this is not a Hess' Law problem.

$$\Delta H = [\Delta H^\circ_{\text{Al}_2(\text{SO}_4)_3(\text{s})} + 6 \Delta H^\circ_{\text{NH}_4\text{NO}_3(\text{s})}] - [2 \Delta H^\circ_{\text{Al}(\text{NO}_3)_3(\text{aq})} + 3 \Delta H^\circ_{(\text{NH}_4)_2\text{SO}_4(\text{aq})}]$$

$$\Delta H = [-820.98 \text{ kcal} + 6(-87.38 \text{ kcal})] - [2(-273.65 \text{ kcal}) + 3(-281.86 \text{ kcal})]$$

$$\Delta H = 47.62 \text{ kcal} \checkmark$$

or

$$199.2 \text{ kJ}$$

4. Determine the final temperature if 70.594 kJ of heat is added to 125 g of ice at 0°C. Useful information can be found in your text on page 307 and of course page 799.

Find heat absorbed by change of state ($\text{H}_2\text{O}(\text{s}) @ 0^\circ\text{C} \rightarrow \text{H}_2\text{O}(\text{l}) @ 0^\circ\text{C}$)

$$Q = L_f m \checkmark$$

$$Q = \left(\frac{6.03 \text{ kJ}}{1 \text{ mol}} \times \frac{1 \text{ mol}}{18.02 \text{ g}} \right) \times 125 \text{ g}$$

$$Q = 41.829 \text{ kJ} \checkmark$$

Find heat not used to cause change of state

$$70.594 \text{ kJ} - 41.829 \text{ kJ} = 28.765 \text{ kJ} \checkmark$$

Use remaining heat to warm water

$$Q = 28.765 \text{ kJ} \rightarrow 28765 \text{ J}$$

$$\Delta T = \frac{Q}{m_c} \checkmark$$

$$m = 125 \text{ g}$$

$$c = 4.184 \text{ J/g}^\circ\text{C}$$

$$\Delta T = ?$$

$$\Delta T = \frac{28765 \text{ J}}{125 \text{ g} \times 4.184 \text{ J/g}^\circ\text{C}}$$

$$\Delta T = 55.00^\circ\text{C} \checkmark$$

∴ the water warms by 55.00°C from 0°C to 55.00°C

5. Determine the mass of propane that must be combusted to warm 160 L of water from 15 °C to 52 °C. You may use any heats of formation from the textbook that you like.

$$Q = ?$$

$$m = 160 \text{ L} \rightarrow 160000 \text{ g}$$

$$c = 4.184 \text{ J/g°C}$$

$$\Delta T = 52^\circ\text{C} - 15^\circ\text{C} = 37^\circ\text{C}$$

$$Q = mc\Delta T \quad \checkmark$$

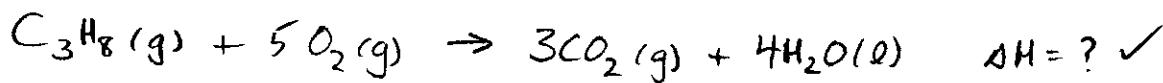
$$Q = 160000 \text{ g} \times 4.184 \text{ J/g°C} \times 37^\circ\text{C}$$

$$Q = 24769280 \text{ J} \quad \checkmark$$

$$Q = 24769 \text{ kJ} \quad \checkmark$$

↑

this is the heat energy required



$$\Delta H = [3\Delta H^\circ_{\text{CO}_2(\text{g})} + 4\Delta H^\circ_{\text{H}_2\text{O}(\text{l})}] - [\Delta H^\circ_{\text{C}_3\text{H}_8(\text{g})} + 5\Delta H^\circ_{\text{O}_2(\text{g})}] \quad \checkmark$$

$$\Delta H = [3(-393.5 \text{ kJ}) + 4(-285.8 \text{ kJ})] - [(-104.7 \text{ kJ}) + 5(0)]$$

$$\Delta H = -2219.0 \text{ kJ} \quad \checkmark$$

$$\Delta H = -Q$$

$$Q = 2219.0 \text{ kJ} / \text{mol C}_3\text{H}_8 \quad \checkmark$$

↑

Heat released per mol of C_3H_8 combustion

$$24769 \text{ kJ} \times \frac{1 \text{ mol C}_3\text{H}_8}{2219.0 \text{ kJ}} \times \frac{44.11 \text{ g C}_3\text{H}_8}{1 \text{ mol C}_3\text{H}_8} = 492.37 \text{ g C}_3\text{H}_8 \quad \checkmark$$

6. If 50 L of water is warmed by 11.469°C by the combustion of 50 g of decane, determine the heat of formation of decane. This is not a Hess' Law problem. You may use any heats of formation that you wish from the textbook.

$$Q = ?$$

$$Q = mc\Delta T \quad \checkmark$$

$$m = 50 \text{ L} \Rightarrow 50000 \text{ g}$$

$$Q = 50000 \text{ g} \times 4.184 \text{ J/g}^{\circ}\text{C} \times 11.469^{\circ}\text{C}$$

$$c = 4.184 \text{ J/g}^{\circ}\text{C}$$

$$Q = 2399314 \text{ J} \quad \checkmark$$

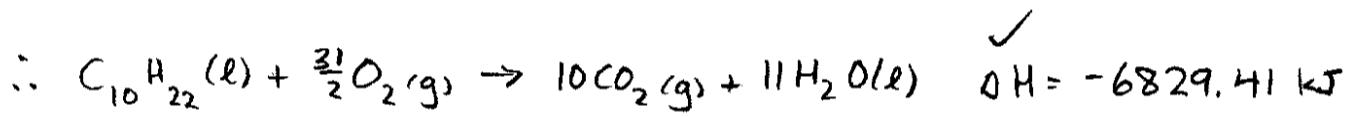
$$\Delta T = 11.469^{\circ}\text{C}$$

$$Q = 2399.31 \text{ kJ} \quad \checkmark$$

$$\frac{2399.31 \text{ kJ}}{50 \text{ g C}_{10}\text{H}_{22}} \times \frac{142.32 \text{ g C}_{10}\text{H}_{22}}{1 \text{ mol C}_{10}\text{H}_{22}} = 6829.41 \text{ kJ/mol C}_{10}\text{H}_{22} \quad \checkmark$$

$$\Delta H = -Q \quad \checkmark$$

$$\Delta H = -6829.41 \text{ kJ/mol C}_{10}\text{H}_{22}$$



$$\Delta H = [10\Delta H^\circ_{\text{CO}_2(g)} + 11\Delta H^\circ_{\text{H}_2\text{O}(l)}] - [\Delta H^\circ_{\text{C}_{10}\text{H}_{22}(l)} + \frac{31}{2}\Delta H^\circ_{\text{O}_2(g)}] \quad \checkmark$$

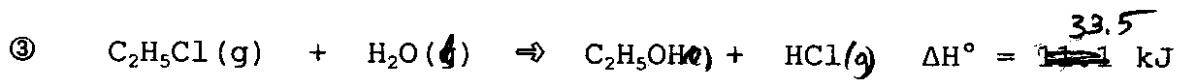
$$-6829.41 \text{ kJ} = [10(-393.5 \text{ kJ}) + 11(-285.8 \text{ kJ})] - [\Delta H^\circ_{\text{C}_{10}\text{H}_{22}(l)} + \frac{31}{2}(0)]$$

$$\therefore \Delta H^\circ_{\text{C}_{10}\text{H}_{22}} = -249.39 \text{ kJ} \quad \checkmark$$

7. Use Hess' Law to find the heat of formation of ethane (C_2H_6) given:



② The heat of combustion of ethanol is -1409.2 kJ .



④ The heat of formation (and reaction thereof) for carbon dioxide gas.

⑤ The heat of formation (and reaction thereof) for water vapour.

⑥ The heat of formation (and reaction thereof) for hydrogen chloride gas.

See text book for heats of formation and the answer!!

